Mage 5 of 5

IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re PATENT APPLICATION OF

Inventor(s): MARTIN ET AL

Filed: Herewith

Title: RELEASE OF INTRACELLULAR MATERIAL

November 28, 2001

PRELIMINARY AMENDMENT

Please amend this application as follows: At the top of the first page, just under the title, insert:		Commissioner of Patents ngton, D.C. 20231
At the top of the first page, just under the title, insert:	C Sig O O E	Please amend this application as follows:
At the top of the first page, just under the title, insert:	INTH	E SPECIFICATION:
Continuation Substitute Application (MPEP 201.09) of 1(a) National Application No. 09/030,028 filed February 25, 1998. Now persent 6335./bl 1(b) International Application No. PCT/GB95/00204 filed August 25, 1995 which designated the U.S 2This application claims the benefit of U.S. Provisional Application No. 60/, filed Respectfully submitted, PILLSBURY WINTHROP LLP Intellectual Property Group Attorney: Paul N. Kokulis Reg. No: 16773 Tel. No.: (703) 905-2118 Fax No.: (703) 905-2500	er E	
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PILLSBURY WINTHROP LLP Intellectual Property Group Attorney: Paul N. Kokulis Reg. No: 16773 Tel. No.: (703) 905-2118 Fax No.: (703) 905-2500		Demostfully submitted
Reg. No: 16773 Tel. No.: (703) 905-2118 Fax No.: (703) 905-2500		PILLSBURY WINTHROP LLP
·		Reg. No: 16773 Tel. No.: (703) 905-2118
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(703) 905-2000

1600 Tysons Boulevard McLean, VA 22102

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IN THE UNITED STATES PATENT AND TRADEMARK OFFICE

In re Application of

MARTIN ET AL

Serial No. Division of 09/030,028

091994 657

Filed: Herewith

RELEASE OF INTRACELLULAR

MATERIAL

Group Art Unit: 1656

Examiner: Tung

November 28, 2001

PRELIMINARY AMENDMENT

Hon. Commissioner of Patents and Trademarks Washington, D.C. 20231

Sir:

For:

Please amend the above divisional application as follows:

IN THE SPECIFICATION

Page 16, 3rd ¶ of Example 3, line 27, change to read as follows:

Two carbon probe electrodes were placed into the sample and 4-8 V (d.c.) was applied (power supply; Thurlby 30V, 2A) for between 0.5 to 2 minutes. The cell debris was pelleted and supernatants were analysed by PCR. PCR conditions were as follows; 0.1 µl/ml of sample in PCR buffer (as above), 1 µM (each) of primers ATGCGTCCGGCCGTAGAGGAT SEQ ID No. 1 and GTATCACGAGGCCCTT SEQ ID No. 2, 200 µM of each of dATP, dCTP, dGTP, dTTP, 5U/ml AmpliTaq DNA polymerase (Perkin Elmer). All reagent concentrations are given as the final concentration in a reaction volume made up with PCR buffer (as above). Amplified

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